This listing of claims will replace all prior versions and listings of claims in the application:

<u>Listing of Claims</u>:

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-29-(Currently amended)

A method for producing an antibody against a Sarcocystis neurona antigen selected from the group consisting of a 16 (± 4) kDa antigen and a 30 (± 4) kDa antigen, as determined by SDS polyacrylamide gel electrophoresis, comprising:

- (a) providing a microorganism containing a DNA encoding a fusion polypeptide in which a Sarcocystis neurona antigen selected from the group consisting of the 16 (±4) kDa antigen and the 30 (±4) kDa antigen is fused to a polypeptide which enables isolation of the fusion polypeptide by affinity chromatography;
- to produce the fusion polypeptide from the DNA;
- (c) isolating the fusion polypeptide from the culture by affinity chromatography;
- isolated by the affinity chromatography antigen with an adjuvant to produce an admixture;

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(e) (c) immunizing a mammal with the admixture containing the fusion polypeptide and the adjuvant to produce antibodies against the 16 kDa antigen or the 30 kDa antigen comprising the fusion polypeptide; and

(f) (d) removing serum from the immunized mammal and isolating from the serum the antibody against the Sarcocystis neurona antigen selected from the group consisting of the (± 4) 16 kDa antigen and the (± 4) 30 kDa antigen.

-30-(Currently amended)

A method for producing a monoclonal antibody against a Sarcocystis neurona antigen selected from the group consisting of a 16 (± 4) kDa antigen and a 30 (± 4) kDa antigen, as determined by SDS polyacrylamide gel electrophoresis, comprising:

- (a) providing a microorganism containing a DNA encoding a fusion polypeptide in which a Sarcocystis neurona antigen selected from the group consisting of the 16 (± 4) kDa antigen and the 30 (± 4) kDa antigen $\pm s$ fused to a polypeptide which enables isolation of the fusion polypeptide by affinity chromatography;
- (b) culturing the microorganism in a culture to produce the fusion polypeptide from the DNA;

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-----(c) isolating the fusion polypeptide from the culture by the affinity chromatography;

(d) (b) admixing the fusion polypeptide isolated by the affinity chromatography antigen with an adjuvant to produce an admixture;

(e) (c) inoculating mice with the admixture containing the fusion polypeptide and the adjuvant to produce antibodies against the 16 kDa antigen or the 30 kDa antigen comprising the fusion polypeptide;

(f) (d) removing the spleens from the mice
which produce the antibodies against the fusion
polypeptide antigen;

(g) (e) removing spleen cells from the spleens and mixing the spleen cells from the spleens with mouse myeloma cells to produce a mixture of fused cells consisting of spleen cells fused to myeloma cells, the spleen cells, and the myeloma cells;

(h) (f) selecting the fused cells on cell culture medium in which the fused cells can grow but in which the spleen cells and the myeloma cells cannot grow; and

(i) (g) screening the fused cells for fused cells which produce the monoclonal antibody against the Sarcocystis neurona antigen selected from the group

consisting of the 16 (± 4) kDa antigen and the 30 (± 4) kDa antigen to produce the monoclonal antibody.

Claims 32-35 (Cancelled)